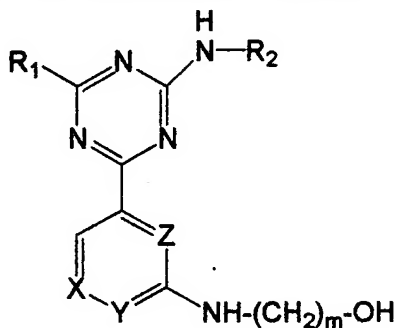


This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

1. (Previously presented) A compound of Formula (I):



Formula (I)

wherein

X, Y and Z are independently selected from the group consisting of CH and N; wherein m is an integer from 2 to 5; wherein X, Y and Z include at least one CH atom and at least one N atom; and, wherein a N atom may simultaneously occupy only the X and Z positions;

R₁ is selected from the group consisting of hydrogen and NH₂; and,

R₂ is selected from the group consisting of phenyl, wherein phenyl is substituted with one substituent selected from the group consisting of halogen and heterocyclyl, and 1,4-benzodioxinyl; or a pharmaceutically acceptable salt thereof.

2. (Original) The compound of claim 1 wherein X, Y and Z are independently selected from the group consisting of CH and N; wherein m is 3; wherein X, Y and Z include at least one CH atom and at least one N atom; wherein a N atom may simultaneously occupy only the X and Z positions; wherein the heteroaryl ring thus formed is selected from the group consisting of pyridinyl and pyrazinyl; wherein pyridinyl is attached to the triazine ring at the 3 or 4 position of the pyridine ring; and, wherein pyrazinyl is attached to the triazine ring at the 6 position of the pyrazine ring;
3. (Original) The compound of claim 1 wherein R₂ is selected from the group consisting of phenyl (wherein phenyl is substituted with one substituent selected from the group consisting of chlorine and 4-morpholinyl) and 1,4-benzodioxinyl.

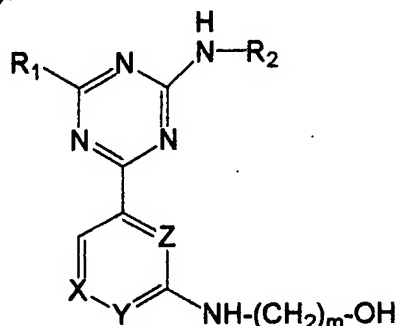
4. (Original) The compound of claim 1 wherein the compound of Formula (I) is selected from a compound wherein m is 3; and, wherein X, Y, Z, R₁ and R₂ are dependently selected from:

X	Y	Z	R ₁	R ₂
N	CH	CH	H	3-Cl-Ph;
CH	N	CH	H	3-Cl-Ph;
N	CH	N	H	3-Cl-Ph;
CH	N	CH	NH ₂	3-Cl-Ph;
N	CH	CH	H	2,3-dihydro-1,4-benzodioxin-6-yl; or,
N	CH	CH	H	4-(4-morpholinyl)Ph.

5. (Original) A composition comprising a compound of claim 1 and a pharmaceutically appropriate carrier.
6. (Canceled)
7. (Original) A method for preparing a composition comprising mixing a compound of claim 1 and a pharmaceutically appropriate carrier.
8. (Canceled)
9. (Canceled)
10. (Canceled)
11. (Canceled)
12. (Canceled)
13. (Canceled)
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18. (Canceled)
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30. (Canceled)
31. (Canceled)
32. (Canceled)
33. (Canceled)
34. (Canceled)
35. (Canceled)
36. (Canceled)

37. (Previously presented) A method for treating human melanoma in a subject in need thereof comprising administering to the subject a therapeutically effective amount of a compound of Formula (I):



Formula (I)

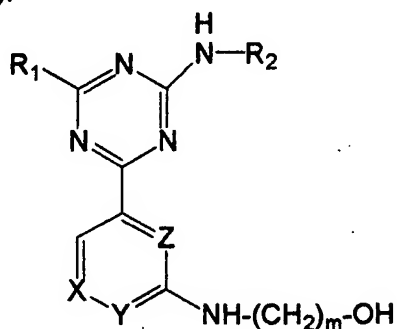
wherein

X, Y and Z are independently selected from the group consisting of CH and N; wherein m is an integer from 2 to 5; wherein X, Y and Z include at least one CH atom and at least one N atom; and, wherein a N atom may simultaneously occupy only the X and Z positions;

R_1 is selected from the group consisting of hydrogen and NH_2 ; and,

R_2 is selected from the group consisting of phenyl, (wherein phenyl is substituted with one substituent selected from the group consisting of halogen and heterocyclyl), and 1,4-benzodioxinyl; or a pharmaceutically acceptable salt thereof.

38. (Previously presented) The method of claim 37 wherein the therapeutically effective amount of the compound of Claim 1 is from about 0.001 mg/kg/day to about 300 mg/kg/day.
39. (Previously presented) A method for treating rheumatoid arthritis in a subject in need thereof comprising administering to the subject a therapeutically effective amount of a compound of Formula (I):



Formula (I)

wherein

X, Y and Z are independently selected from the group consisting of CH and N; wherein m is an integer from 2 to 5; wherein X, Y and Z include at least one CH atom and at least one N atom; and, wherein a N atom may simultaneously occupy only the X and Z positions;

R1 is selected from the group consisting of hydrogen and NH₂; and,

R2 is selected from the group consisting of phenyl, (wherein phenyl is substituted with one substituent selected from the group consisting of halogen and heterocyclyl), and 1,4-benzodioxinyl;

or a pharmaceutically acceptable salt thereof.

40. (Previously presented) The method of claim 39 wherein the therapeutically effective amount of the compound of Claim 1 is from about 0.001 mg/kg/day to about 300 mg/kg/day.